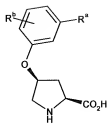


Amendments to the Claims:

1. - 10. (Canceled)
11. (Currently Amended) A pharmaceutical composition comprising a compound of formula ~~(I)~~(Ia), ~~as described in according to claim 14, or a pharmaceutically acceptable salt thereof,~~ and one or more pharmaceutically acceptable excipients, diluents or carriers.
12. (Currently Amended) A combination comprising a compound of formula ~~(I)~~(Ia), ~~as described in according to claim 14, or a pharmaceutically acceptable salt, solvate or pre-drug thereof, and at least one other therapeutically active agent.~~
13. (Original) A combination according to claim 12, wherein the other therapeutically active agent is a PDEV inhibitor.

14. (New) A compound of formula (Ia):



(Ia)

wherein R^a is selected from halogen, hydroxy, $(\text{C}_1\text{-C}_6)$ alkoxy, cyano, nitro, amino, hydroxycarbonyl,

$\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkenyl, $\text{C}_1\text{-C}_6$ alkynyl,

hydroxy $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkoxy $\text{C}_1\text{-C}_6$ alkyl, perfluoro $\text{C}_1\text{-C}_6$ alkyl, perfluoro $\text{C}_1\text{-C}_6$ alkoxy,

$\text{C}_1\text{-C}_6$ alkylamino, di- $\text{C}_1\text{-C}_6$ alkylamino, amino $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkylamino $\text{C}_1\text{-C}_6$ alkyl, di- $\text{C}_1\text{-C}_6$ alkylamino $\text{C}_1\text{-C}_6$ alkyl,

$\text{C}_1\text{-C}_6$ acyl, $\text{C}_1\text{-C}_6$ acyloxy, $\text{C}_1\text{-C}_6$ acyloxy $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ acylamino,

$\text{C}_1\text{-C}_6$ alkylthiocarbonyl, $\text{C}_1\text{-C}_6$ alkylthio, $\text{C}_1\text{-C}_6$ alkoxycarbonyl,

$\text{C}_1\text{-C}_6$ alkylsulfonyl, $\text{C}_1\text{-C}_6$ alkylsulfonylamino,

aminosulfonyl, $\text{C}_1\text{-C}_6$ alkylaminosulfonyl, di- $\text{C}_1\text{-C}_6$ alkylaminosulfonyl,

3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl;

R^b is selected from hydrogen, halogen, hydroxy, $(\text{C}_1\text{-C}_6)$ alkoxy cyano, nitro, amino, hydroxycarbonyl,

$\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkenyl, $\text{C}_1\text{-C}_6$ alkynyl,

hydroxy $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkoxy $\text{C}_1\text{-C}_6$ alkyl, perfluoro $\text{C}_1\text{-C}_6$ alkyl, perfluoro $\text{C}_1\text{-C}_6$ alkoxy,

$\text{C}_1\text{-C}_6$ alkylamino, di- $\text{C}_1\text{-C}_6$ alkylamino, amino $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ alkylamino $\text{C}_1\text{-C}_6$ alkyl, di- $\text{C}_1\text{-C}_6$ alkylamino $\text{C}_1\text{-C}_6$ alkyl,

$\text{C}_1\text{-C}_6$ acyl, $\text{C}_1\text{-C}_6$ acyloxy, $\text{C}_1\text{-C}_6$ acyloxy $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_1\text{-C}_6$ acylamino,

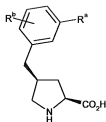
$\text{C}_1\text{-C}_6$ alkylthio, $\text{C}_1\text{-C}_6$ alkylthiocarbonyl, $\text{C}_1\text{-C}_6$ alkylthio, $\text{C}_1\text{-C}_6$ alkoxycarbonyl,

$\text{C}_1\text{-C}_6$ alkylsulfonyl, $\text{C}_1\text{-C}_6$ alkylsulfonylamino,

aminosulfonyl, $\text{C}_1\text{-C}_6$ alkylaminosulfonyl, di- $\text{C}_1\text{-C}_6$ alkylaminosulfonyl,

3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

15. (New) A compound of formula (Ib):



(Ib)

wherein R^a is selected from halogen, hydroxy, cyano, nitro, amino, hydroxycarbonyl, C_1 - C_6 alkyl, C_1 - C_6 alkenyl, C_1 - C_6 alkynyl, hydroxy C_1 - C_6 alkyl, C_1 - C_6 alkoxy C_1 - C_6 alkyl, perfluoro C_1 - C_6 alkoxy, C_1 - C_6 alkylamino, di- C_1 - C_6 alkylamino, amino C_1 - C_6 alkyl, C_1 - C_6 alkylamino C_1 - C_6 alkyl, di- C_1 - C_6 alkylamino C_1 - C_6 alkyl,

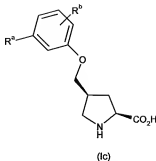
C_1 - C_6 acyl, C_1 - C_6 acyloxy, C_1 - C_6 acyloxy C_1 - C_6 alkyl, C_1 - C_6 acylamino, C_1 - C_6 alkylthio, C_1 - C_6 alkylthiocarbonyl, C_1 - C_6 alkylthio, C_1 - C_6 alkoxycarbonyl, C_1 - C_6 alkylsulfonyl, C_1 - C_6 alkylsulfonylamino, aminosulfonyl, C_1 - C_6 alkylaminosulfonyl, di- C_1 - C_6 alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl;

R^b is selected from hydrogen, halogen, hydroxy, (C_1 - C_6)alkoxy cyano, nitro, amino, hydroxycarbonyl,

C_1 - C_6 alkyl, C_1 - C_6 alkenyl, C_1 - C_6 alkynyl, hydroxy C_1 - C_6 alkyl, C_1 - C_6 alkoxy C_1 - C_6 alkyl, perfluoro C_1 - C_6 alkyl, perfluoro C_1 - C_6 alkoxy, C_1 - C_6 alkylamino, di- C_1 - C_6 alkylamino, amino C_1 - C_6 alkyl, C_1 - C_6 alkylamino C_1 - C_6 alkyl, di- C_1 - C_6 alkylamino C_1 - C_6 alkyl,

C_1 - C_6 acyl, C_1 - C_6 acyloxy, C_1 - C_6 acyloxy C_1 - C_6 alkyl, C_1 - C_6 acylamino, C_1 - C_6 alkylthio, C_1 - C_6 alkylthiocarbonyl, C_1 - C_6 alkylthio, C_1 - C_6 alkoxycarbonyl, C_1 - C_6 alkylsulfonyl, C_1 - C_6 alkylsulfonylamino, aminosulfonyl, C_1 - C_6 alkylaminosulfonyl, di- C_1 - C_6 alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

16. (New) A compound of formula (1c):



wherein R^a and R^b are independently selected from hydrogen, halogen, hydroxy, (C_1-C_6) alkoxy, cyano, nitro, amino, hydroxycarbonyl, C_1-C_6 alkyl, C_1-C_6 alkenyl, C_1-C_6 alkynyl, hydroxy C_1-C_6 alkyl, C_1-C_6 alkoxy C_1-C_6 alkyl, perfluoro C_1-C_6 alkyl, perfluoro C_1-C_6 alkoxy, C_1-C_6 alkylamino, di- C_1-C_6 alkylamino, amino C_1-C_6 alkyl, C_1-C_6 alkylamino C_1-C_6 alkyl, di- C_1-C_6 alkylamino C_1-C_6 alkyl, C_1-C_6 acyl, C_1-C_6 acyloxy, C_1-C_6 acyloxy C_1-C_6 alkyl, C_1-C_6 acylamino, C_1-C_6 alkylthio, C_1-C_6 alkylthiocarbonyl, C_1-C_6 alkylthio, C_1-C_6 alkoxycarbonyl, C_1-C_6 alkylsulfonyl, C_1-C_6 alkylsulfonylamino, aminosulfonyl, C_1-C_6 alkylaminosulfonyl, di- C_1-C_6 alkylaminosulfonyl, 3-8 membered cycloalkyl, 4-8 membered heterocycloalkyl, phenyl and monocyclic heteroaryl; or a pharmaceutically acceptable salt thereof.

17. (New) A compound of formula (1a) according to claim 1 which is:

(2S, 4S)-4-(3-Chloro-phenoxy)-pyrrolidine-2-carboxylic acid;

or a pharmaceutically acceptable salt thereof.

18. (New) A compound of formula (1b) according to claim 15 which is selected from the group consisting of:

(2S,4S)-4-(3-Fluoro-benzyl)-pyrrolidine-2-carboxylic acid;

(2S,4S)-4-(2,3-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; and

(2S,4S)-4-(2,5-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; or a pharmaceutically acceptable salt thereof.

19. (New) A compound of formula (1c) according to claim 16 which is selected from the group consisting of:

(2S,4S)-4-(3-Fluoro-phenoxy-methyl)-pyrrolidine-2-carboxylic acid;

(2S,4S)-4-(3,6-Difluoro-phenoxyethyl)-pyrrolidine-2-carboxylic acid;
(2S,4S)-4-(2,3-Difluoro-phenoxyethyl)-pyrrolidine-2-carboxylic acid; and
(2S,4S)-4-(3-Methoxy-phenoxyethyl)-pyrrolidine-2-carboxylic acid; or a pharmaceutically acceptable salt thereof.

20. (New) A pharmaceutical composition comprising a compound of formula (Ib) according to claim 15, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.

21. (New) A combination comprising a compound of formula (Ib) according to claim 15, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.

22. (New) A combination according to claim 21, wherein the other therapeutically active agent is a PDEV inhibitor.

23. (New) A pharmaceutical composition comprising a compound of formula (Ic) according to claim 16, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.

24. (New) A combination comprising a compound of formula (Ic) according to claim 16, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.

25. (New) A combination according to claim 24, wherein the other therapeutically active agent is a PDEV inhibitor.

26. (New) The compound (2S,4S)-4-(3-Chloro-phenoxy)-pyrrolidine-2-carboxylic acid or a pharmaceutically acceptable salt thereof.